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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/547,995	. 10/28/2005	David John Grainger	50461/003001	8214
21559 CLARK & ELI	7590 12/31/2007 BING LLP		EXAM	INER
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BOSTON, MA 02110		ART UNIT	PAPER NUMBER	
			1641	
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			NOTIFICATION DATE	DELIVERY MODE
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

patentadministrator@clarkelbing.com

	Application No.	Applicant(s)			
•	10/547,995	GRAINGER, DAVID JOHN			
Office Action Summary	Examiner	Art Unit			
	Christine Foster	1641			
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply					
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA. - Extensions of time may be available under the provisions of 37 CFR 1.1: after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period variety or Failure to reply within the set or extended period for reply will, by statute, any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNIONS 36(a). In no event, however, may a rickly and will expire SIX (6) MON, cause the application to become AB	CATION. eply be timely filed THS from the mailing date of this communication. ANDONED (35 U.S.C. § 133).			
Status					
 1) ⊠ Responsive to communication(s) filed on 15 Dec 2a) ☐ This action is FINAL. 2b) ☒ This 3) ☐ Since this application is in condition for allower closed in accordance with the practice under E 	action is non-final.	•			
Disposition of Claims					
4) Claim(s) <u>1-46</u> is/are pending in the application. 4a) Of the above claim(s) is/are withdray 5) Claim(s) is/are allowed. 6) Claim(s) is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) <u>1-46</u> are subject to restriction and/or expressions.	vn from consideration.				
Application Papers					
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) acce Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex	epted or b) objected to drawing(s) be held in abeyand ion is required if the drawing	ice. See 37 CFR 1.85(a). (s) is objected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119					
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 					
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	Paper No(s	Summary (PTO-413) s)/Mail Date nformal Patent Application			

10/547,995 Art Unit: 1641

DETAILED ACTION

Election/Restrictions

1. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

- Group I, claim(s) 1-12 and 46, drawn to a method for determining the relative abundance of a plurality of proteins.
- Group II, claim(s) 13-21 and 41-44, drawn to peptide mixtures and libraries and kits comprising same.
- Group III, claim(s) 22-33, drawn to a method of detecting a plurality of immunoglobulins.
- Group IV, claim(s) 34-40, drawn to a method of detecting the presence of, or a susceptibility to, a disease or other medical condition.
- Group V, claim(s) 45, drawn to a method of reducing the redundancy and bias of an antibody-expressing phage library.
- 2. The inventions listed as Groups I-V do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

According to PCT Rule 13.2, unity of invention exists only when the shared same or corresponding technical feature is a contribution over the prior art. The technical feature of Group I is that of a method of determining the relative abundance of a plurality of proteins using

10/547,995 Art Unit: 1641

as reagents (1) a reference sample comprising labeled proteins and (2) a plurality of tagged antibodies.

Chandler et al. (US 6,449,562 B1) teach methods of determining the levels of multiple analytes in fluid samples using bead-based flow cytometry methods. In one embodiment, a plurality of proteins (IgG, IgA, IgM, and BSA) was labeled by coating with fluorescent microspheres in order to simultaneously determine the levels of IgG, IgA, and IgM in a test serum sample (column 23, line 66 to column 25, line 45). The Ig-coated microspheres are incubated with the test sample (calibrator sera serving as unknowns) in the presence of a plurality of tagged antibodies (mixture of three Bodipy-labeled anti-human Ig antibodies) (see in particular column 25, lines 27-44 and column 27, lines 6-43). The amount of fluorescence Fm produced by the Bodipy-labeled antibodies binding to the microspheres (via sample antibody bridge) is then detected (see also column 23, lines 11-19; column 24, lines 14-27; and Tables 3-5). The results are compared to those of serum calibrators containing known amounts of IgG, IgA, and IgM (column 25, lines 27-44; column 27, lines 7-16), which may be zero (Tables 3-5), i.e. no test sample. Therefore, in light of the teachings of Chandler et al., the technical feature of Group I does not constitute a special technical feature as defined by PCT Rule 13.2, as it does not define a contribution over the prior art.

Election of Species

3. This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

Art Unit: 1641

The species are as follows:

In the event that <u>Group I</u> above is elected, the following elections are also required:

- a. Antibody tag linkage (elect one of the following):
 - i. Each tag is linked to a single antibody species (see claim 4).
 - ii. Each tag is linked to more than one species of antibody (see claim 5)
- b. Antibody binding (elect one of the following):
 - i. Each antibody species binds the same protein (see claim 6)
 - ii. Each tagged antibody binds a different protein (see claim 7)

Applicant is required, in reply to this action, to elect a single species to which the claims shall be restricted if no generic claim is finally held to be allowable. The reply must also identify the claims readable on the elected species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

The claims are deemed to correspond to the species listed above as indicated. The following claim(s) are generic: claims 1-3, 8-12, and 46 appear to be generic.

10/547,995

Art Unit: 1641

4. The species listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons:

Chandler et al. (discussed above) exemplifies adding a mixture of three anti-Ig antibodies (anti-IgG, anti-IgA, and anti-IgM), each of which are labeled with the Bodipy tag (column 27, lines 13-16), such that the tag is linked to more than one species or type of antibody. These anti-Ig antibodies bind to different proteins (i.e., IgG, IgA, and IgM immunoglobulin, respectively). Therefore, the species indicated above are not regarded as being of similar nature because they are not linked by a common property or activity that defines a contribution over the prior art.

Applicant is advised that the reply to this requirement to be complete must include (i) an election of a species or invention to be examined even though the requirement be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.

The election of an invention or species may be made with or without traverse. To reserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse.

Should applicant traverse on the ground that the inventions or species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the inventions or species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C.103(a) of the other invention.

10/547,995 Art Unit: 1641

Notice of Possible Rejoinder

The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and the product claims are subsequently found allowable, withdrawn process claims that depend from or otherwise require all the limitations of the allowable product claim will be considered for rejoinder. All claims directed a nonelected process invention must require all the limitations of an allowable product claim for that process invention to be rejoined.

In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103 and 112. Until all claims to the elected product are found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowable product claim will not be rejoined. See MPEP § 821.04(b). Additionally, in order to retain the right to rejoinder in accordance with the above policy, applicant is advised that the process claims should be amended during prosecution to require the limitations of the product claims. Failure to do so may result in a loss of the right to rejoinder. Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Christine Foster whose telephone number is (571) 272-8786. The examiner can normally be reached on M-F 8:30-5. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached at (571) 272-0823. The fax

Application/Control Number:

10/547,995 Art Unit: 1641 Page 7

phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Chost

Christine Foster
Patent Examiner
Art Unit 1641

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SUPERVISORY PATENT EXAMINER
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